

## **REMARKS**

### **Status of Claims**

Claims 1, 3-25 and 33-38 are pending. Claim 1 has been amended. The amendment does not include new matter and is fully supported by the application. In particular, support for amended claim 1 can be found, *inter alia*, on pages 11-12 of the application, as originally filed.

### **Applicants' Response to 35 U.S.C. §102 Rejection over Short as evidenced by Alberts**

Claims 1, 3-13, 15-16 and 36 were rejected under 35 U.S.C. §102(b) as being allegedly anticipated by WO Publication No. WO 01/31339 to Short et al. (hereinafter "Short") as evidenced by Molecular Biology of the Cell to Alberts et al. (hereinafter "Alberts").

The Examiner alleged that:

Short teaches the steps, as part of said biological assay, in which a biological molecule is bound to a surface which has been treated by plasma polymerization. The step of 'binding' is reasonably interpreted as the biological molecule *contacting* the plasma treated surface. It is further the position of the Examiner, absent a showing of evidence to the contrary, that the newly added limitation of passively absorbed [sic] on the surface 'and thereby immobilized' recited in instant Claim 1, step v, is met by the disclosure of Short that the biological molecule is bound...the passively absorbed [sic] and thereby immobilized carbohydrate molecule retains its biological activity, is accorded no patentable weight as such an outcome would necessarily result from the method steps taught by Short which, as noted above, are the same as those recited in the instant Claim.

(Office Action, at page 4) (citations omitted).<sup>1</sup>

In the Advisory Action, the Examiner referred to the arguments made in Applicants' June 9, 2010 response. In particular, the Examiner stated that the arguments were not persuasive because "Applicant has not provided a limiting definition of 'contaminated', merely a preference

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<sup>1</sup> In the Advisory Action dated June 28, 2010, the Examiner acknowledged that references to "absorption" were incorrect and that "Clearly, the intent was to indicate that the substrates of Short are 'adsorbed', i.e. bound to the surface by an attractive force." Accordingly, Applicants will respond under the assumption that the Examiner's use of "absorbed" in the Final Office Action was meant to be "adsorbed" in the following response.

that, in assays, the polysaccharide is adsorbed in the absence of albumin or salts.” Claim 1 has been amended to indicate that the passive adsorption is done “in the absence of albumin or salt” and that the carbohydrates are “not contaminated”.

Short discloses plasma polymerization of surfaces to which antagonists may bind and be assayed for use in the detection and/or activity of a biological molecule bound thereto. Nowhere in Short is it disclosed or suggested to provide a non-contaminated carbohydrate, in its native form, and passively adsorb the non-contaminated carbohydrate in its native form onto a plasma polymerized surface, in the absence of albumin or salts.

The Examiner stated on page 14 of the Office Action that “the antibody of Short, human immunoglobulin G (IgG), appears, absent any specific evidence to the contrary to contact directly the plasma polymer coated surface without prior modification of the coated surface.” As set forth at page 12, ll. 7-9, the IgG is diluted in phosphate buffered saline (PBS) and then exposed to a surface. As set forth above, with the method of claim 1, the carbohydrate molecule is passively adsorbed in the absence of salts. The IgG in Short is adsorbed in the presence of salt.

It is respectfully submitted that claims 1, 3-13, 15-16, 36 and 38 are patentable over Short.

**Applicants' Response to 35 U.S.C. §102 Rejection over Yan**

Claims 1, 3, 5-7, 11, 18-20 and 22 were rejected under 35 U.S.C. §102(b) as being allegedly anticipated by U.S. Patent No. 6,776,792 to Yan et al. (hereinafter “Yan”).

The Examiner alleged that:

Yan et al. teach an implantable stent coated with a material that attracts the glycosaminoglycan carbohydrate heparin and forms a bond. Yan teaches that the stent surface coating is formed by plasma deposition with methane, ammonia gas,

other amine containing monomers and acrylic acid. Yan also teaches contacting the plasma coated surface with heparin, via ionic bonds. Finally, Yan teaches the incubation step recited in amended Claim 1 by pretreating the plasma coated surface (stent) with a heparinized saline solution prior to implantation in order to adjust the heparin level. Therefore, the carbohydrate is immobilized using the method steps recited in instant Claim 1 and accordingly, such an outcome results in heparin being passively (ionically) absorbed and retaining its anticoagulant biological activity.

(Office Action, at page 9) (citations omitted)

Yan is directed to a coated endovascular stent. The stent is coated with a material that attracts heparin so that the heparin thus forms a bond with the material. (See, e.g., col. 2, ll. 57-60). The purpose of Yan is to prepare a stent that is biocompatible upon implantation. Yan includes a coating with functional groups that attract heparin. Yan describes a plasma deposition as a possible embodiment to obtain the coating. The steps for the plasma deposition include depositing a base or primer layer to "prepare the surface of the stent to receive the functionality group containing substance" (Yan, col. 3, ll. 4-5) and a second or "top layer that includes the desired functionalities" (Yan, col. 3, ll. 9-10). Accordingly, one of skill in the art would appreciate that the molecules of Yan are not passively adsorbed onto the surface in its native state.

On page 15 of the Office Action, the Examiner pointed to the discussion in Yan regarding a physician pre-treating the stent with a heparin saline solution in arriving at the conclusion that the molecules must be immobilized. As stated above, amended claim 1 requires that the carbohydrate be passively adsorbed without utilizing salts or albumin. In contrast, Yan utilizes a salt.

It is respectfully submitted that claims 1, 3, 5-7, 11, 18-20 and 22 are patentable over Yan.

**Applicants' Response to 35 U.S.C. §103 Rejection over Short as evidenced by Alberts and Karwoski**

Claims 14 and 33-34 were rejected under 35 U.S.C. §103(a) as being allegedly unpatentable over Short as evidenced by Alberts and further in view of U.S. Patent No. 4,632,842 to Karwoski et al. (hereinafter "Karwoski").

As stated above, Short fails to disclose each and every element of claim 1. The Examiner cited Karwoski for disclosing the formula W/FM and alleged that "one would have been motivated to use the ranges for W/FM...because these ranges have already been successfully used by Short to provide plasma coated assay binding surfaces containing the same genus of plasma monomers." (Office Action, at page 8). Karwoski is directed to a glow discharge process for producing implantable devices. Nowhere in Karwoski is it disclosed or suggested to utilize a non-contaminated carbohydrate in its native form or a polymerized surface that has not been modified prior to contact with the carbohydrate in its native form. Accordingly, Karwoski fails to cure the deficiencies of Short.

It is respectfully submitted that claims 14 and 33-34 are patentable over Short, Alberts and Karwoski, each taken alone or in combination.

**Applicants' Response to 35 U.S.C. §103 Rejection over Yan, and further in view of Mori**

Claim 17 was rejected under 35 U.S.C. §103(a) as being allegedly unpatentable over Yan in view of U.S. Patent No. 5,053,398 to Mori et al. (hereinafter "Mori").

As stated above, Yan fails to disclose each and every element of claim 1. Mori discloses sulfated homopolysaccharides with anti-HIV activity. However, Mori does not disclose or suggest immobilization of such homopolysaccharides. Furthermore, Mori fails to disclose or suggest passively adsorbing non-contaminated homopolysaccharides without utilizing salt or albumin, and, as such, Mori fails to overcome the deficiencies of Yan in this regard.

It is respectfully submitted that claim 17 is patentable over Yan and Mori, each taken alone or in combination.

**Applicants' Response to 35 U.S.C. §103 Rejection over Short as evidenced by Alberts and Nomura**

Claim 35 was rejected under 35 U.S.C. §103(a) as being allegedly unpatentable over Short as evidenced by Alberts and further in view of U.S. Patent No. 6,022,602 to Nomura et al. (hereinafter "Nomura").

As stated above, Short fails to disclose each and every element of claim 1. Nomura is directed to a lumen surface of tubing that has been modified with plasma. Nomura was merely cited for its disclosure of an alkane. Accordingly, Nomura fails to cure the deficiencies of Short.

It is respectfully submitted that claim 35 is patentable over Short, Alberts and Nomura, each taken alone or in combination.

**Applicants' Response to 35 U.S.C. §103 Rejection over Short, and further in view of Nilsson**

Claim 21 was rejected under 35 U.S.C. §103(a) as being allegedly unpatentable over Short and further in view of U.S. Publication No. 2001/0017270 to Nilsson et al. (hereinafter "Nilsson").

As stated above, Short fails to disclose each and every element of claim 1, as amended herein. Nilsson refers to a biosensor in which a carbohydrate or a derivative thereof is bound to a surface of the biosensor *via* a chemical bond or adsorption. In particular, Nilsson states "the surface of the biosensor can be, for example a gold surface or a modified gold surface, a plastic surface which has been modified with a gold surface, silver surface or another metallic surface, or modifications thereof with polymers to which **chemical coupling** of carbohydrate can be

carried out." (See Nilsson, page 3, paragraph [0041], emphasis added). However, nowhere does Nilsson describe any modifications to a plasma polymer surface for passive adsorption of a carbohydrate. Likewise, nowhere does Nilsson exemplify a biosensor with a polymer surface. Rather, Nilsson exemplifies the following:

- a) binding of digalactoside with aglycon (*i.e.*, Gal $\alpha$ 1-4Gal $\beta$ -OEtSeTCONHNH<sub>2</sub>) to a silica surface coated with a gold layer modified with mercaptopropionic acid
- b) binding of Gal $\alpha$ 1-4Gal $\beta$ OCH<sub>2</sub>CH<sub>2</sub>SCH<sub>2</sub>C(O)-NHNH -BSA to a silica surface coated with a gold layer modified with mercaptopropionic acid
- c) binding of Gal $\alpha$ 1-4Gal $\beta$ -BSA to a silica surface coated with a gold layer (not pretreated with mercaptopropionic acid)

Thus, Nilsson teaches biosensors with a surface coated with a gold layer. Nowhere in Nilsson is it disclosed or suggested to provide a plasma polymerized surface that is not modified prior to coming into contact with a carbohydrate. Moreover, nowhere in Nilsson is utilizing a carbohydrate in its native form disclosed or suggested. Accordingly, Nilsson does not overcome the deficiencies noted above of Short.

It is respectfully submitted that claim 21 is patentable over Short, Alberts and Nilsson, each taken alone or in combination.

**Applicants' Response to 35 U.S.C. §103 Rejection over Yan, and further in view of Earhart**

Claim 23 was rejected under 35. U.S.C. §103(a) as being allegedly unpatentable over Yan in view of U.S. Patent No. 6,077,232 to Earhart et al. (hereinafter "Earhart").

As stated above, Yan fails to disclose each and every element of claim 1, as amended herein. Earhart teaches a blood collection device containing an anticoagulant composition. Specifically, Earhart states that,

The anticoagulant composition is a soluvated non-aqueous solution comprising a proteinase inhibitor such as for example but not limited to a polysaccharide sulfuric acid ester such as for example heparin. The composition also comprises a blend of one or more alcohols such as for example one or more polyhydric alcohols selected from the group consisting of C1-12 diols such as glycols and derivatives thereof such as for example ethylene glycol or propylene glycol wherein ethylene glycol is preferred for increased solubility and C1-12 polyalkyl diols such as for example polyethylene glycol, polypropylene glycol or polybutylene glycol wherein polyethylene glycol is preferred for increased solubility and for ready plasticization of the proteinase inhibitor.

(col. 2, l. 62 – col. 3, l.8).

Furthermore, Earhart states that “the subject anticoagulant composition is placed within the syringe barrel in a liquid state...and remains in the syringe barrel during blood collection due to its movement within the barrel without expulsion thereof.” (col. 1, l. 62 to col. 2, l. 6). Thus, the anticoagulant composition is not immobilized to the surface but rather in a liquid state that freely moves. In fact, Earhart further emphasizes that, “The alcohol which is preferably polyethylene glycol, combined with the proteinase inhibitor which is preferably heparin, prevents the proteinase inhibitor from drying into an unsolvated state. Heparin in an unsolvated state dissolves with difficulty and thus requires more time to disperse throughout a collected blood sample.” (col. 2, ll. 15-21). In fact, Earhart points out that “the use of the anticoagulant composition of the present invention within a blood collection device reduces the anticoagulant solution dispersal time compared to that of dried unsolvated heparin within a blood collection device so as to be more effective in preventing coagulation within a collected blood sample.” Thus, the proteinase inhibitor, heparin, is preferably a liquid which disperses readily throughout a blood sample collected in the blood collection device. Nowhere does Earhart disclose or suggest immobilizing heparin. Without immobilization, it is respectfully submitted that Earhart does not overcome the deficiencies noted above of Yan.

It is respectfully submitted that claim 23 is patentable over Yan and Earhart, each taken alone or in combination.

**Applicants' Response to 35 U.S.C. §103 Rejection over Short, and further in view of Brigstock**

Claim 24 was rejected under 35 U.S.C. §103(a) as being allegedly unpatentable over Short and further in view of U.S. Publication No. 2001/0007019 to Brigstock et al. (hereinafter "Brigstock").

As stated above, Short fails to disclose each and every element of claim 1, as amended herein. Brigstock teaches heparin-binding growth factor (HBGF) polypeptides as well as nucleic acids encoding the same and antibodies which bind to the HBGF polypeptides. Brigstock discloses the use of a heparin affinity chromatographic column in its purification of HBGF polypeptides. (page 7, paragraphs [0060] to [0063]). However, no additional detail regarding such a heparin affinity chromatographic column is provided. With no details on whether or not the polypeptides are immobilized, Brigstock does not overcome the deficiencies noted above of Short.

It is respectfully submitted that claim 24 is patentable over Short and Brigstock, each taken alone or in combination.

**Applicants' Response to 35 U.S.C. §103 Rejection over Short, and further in view of Dukler**

Claim 25 was rejected under 35 U.S.C. §103(a) as being allegedly unpatentable over Short and further in view of U.S. Publication No. 2002/0094541 to Dukler et al. (hereinafter "Dukler").

As stated above, Short fails to disclose each and every element of claim 1, as amended herein. Dukler was cited for its alleged disclosure of a microarray. Dukler teaches combinatorial complex libraries and methods for the manufacture of addressable complex carbohydrate microarrays as well as uses thereof. As the carbohydrate microarrays of Dukler are addressable, one of skill in the art would not be motivated to employ passive adsorption of

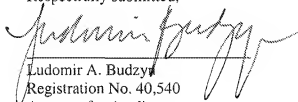
carbohydrates to the substrate surface since such non-covalent bonds would be impractical to produce in an addressable format. Dukler states that, "the libraries according to the present invention are preferably synthesized on a solid phase support. As such, the first building block is provided with a suitable functional group for binding such a support." (page 15, paragraph [0165]). Furthermore, with respect to linking the first saccharide building block to the solid phase support, Dukler states, "The first saccharide building block is preferably covalently attached to the solid phase matrix via a single atom (e.g., the solid phase functional group) or a linker." (page 15, paragraph [0154]). In fact, Dukler states, "... covalent immobilization methods enable the use of a very high ionic strength buffer (e.g., 6 M Guanidine HCl or 100 mM NaOH) in subsequent washing steps thus allowing accurate "in situ" verification of each enzymatic step utilized by the process. The removal of nonspecifically bound molecules is crucial for accurate library synthesis." (page 49, paragraph [0242] and [0243], first sentence). Thus, one of skill in the art reading Dukler would choose covalent attachment over non-covalent attachment of a carbohydrate to a surface of the microarray. Nowhere in Dukler is it disclosed or suggested to utilize a carbohydrate in its native form or a polymerized surface that has not been modified prior to contact with the carbohydrate in its native form. Accordingly, Dukler does not overcome the deficiencies noted above of Short.

It is respectfully submitted that claim 25 is patentable over Short and Dukler, each taken alone or in combination.

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Favorable action is earnestly solicited. If there are any questions or if additional information is required, the Examiner is respectfully requested to contact Applicants' attorney at the number listed below.

Respectfully submitted,



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